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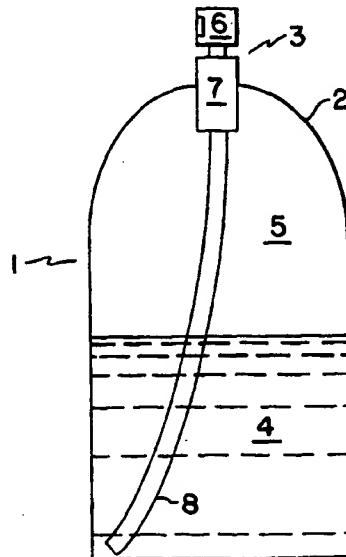
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(54) Title: DISPENSING DISPERSED POWDER WHICH RESOLUBILIZES ON APPLICATION

(57) Abstract

The dispensing system (1) includes a container (2). Container (2) is a sealed enclosure to which is fitted externally actuable dispensing means (3) for dispensing product when desired from within container (2) to the ambient atmosphere or to a target surface. Within container (2) are a dispersion (4) of active solids in a homogeneous mixture of solvent and propellant (5). Dispensing means (3) includes a button (6), valve means (7), and dip tube (8). Valve means (7) is preferably of the type having a passage for product being dispensed, and a valve operatively coupled to a spring means which urges the valve to a normally closed position preventing passage of product therethrough except when the valve is urged to an opening position by external actuation.



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DISPENSING DISPERSED POWDER WHICH
RESOLUBILIZES ON APPLICATION

1

The present invention relates to dispensing systems. It relates more particularly to dispensing systems in which a sealed container contains a product of an insoluble active ingredient and solvent therefor all mixed with a volatile propellant or liquid. In a suitable dispensing system, the product is released, typically in the form of a stream, mist, foam, or fog of droplets having active ingredient redissolved in the solvent.

Dispensers of this type have often been employed for dispensing liquid solutions. Typically, the container holds the solution under pressure exerted by the propellant. When the solution is dispensed from the container upon actuation of the valve means, it remains in solution as it emerges from the container and is applied to a target surface as a solution. In addition, pressurized dispensing systems have also been used for dispensing finely divided solids. The solids are held within the container dispersed in liquefied propellant and/or in a liquid carrier. When the dispersion is dispensed from the container upon actuation of the valve means, the solids emerge for application to a target surface. Examples of products which are dispensed in this manner include antiperspirants, paints and other surface treatments, lubricants, pesticides, and so forth.

The prior art relating to aerosol dispensing of antiperspirants illustrates the manner in which pressurized dispensers have heretofore been used to dispense solutions and dispersions of solids.

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1 Antiperspirants have classically presented a challenge
to dispense because they can be difficult to solubilize.

Thus, the approaches heretofore taken to
formulating systems for dispensing active ingredients
5 such as antiperspirants from pressurized, aerosol-type
containers, have followed two courses. One course has
been to formulate the product into a solution containing
the active ingredient, whereby the formulation is in
solution form within the pressure container, as
10 discharged from the container, and as applied to a
target surface. The other course has been to formulate
the active ingredient into an emulsion or a dispersion
of the active ingredient as a solid phase, wherein the
active ingredient is in solid form within the container,
15 and remains in solid form as the emulsion or dispersion
is discharged, and upon application to a target surface.

For instance, U.S. Patent No. 3,928,545, U.S.
Patent No. 3,947,556, and U.S. Patent No. 3,904,741
disclose alcohol-soluble complexes of basic aluminum
20 chlorides with zirconyl or zinc compounds, which are
said to be useful in preparing aerosol antiperspirant
sprays. These patents have as their objective the
formation of solutions characterized in that the
antiperspirant active ingredient remains solubilized in
25 the container and upon discharge therefrom. These
patents speak of the solutions having good "fluorocarbon
compatibility" which refers to the ability of the
solution to retain all of the active ingredient in
solution even in the presence of fluorocarbon
30 propellants within the pressurized container, such that

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1 the active ingredient does not precipitate from solution
as a solid.

Other patents describe similarly alcoholic or
hydro-alcoholic solutions of antiperspirants, and add
5 that having the active ingredient take solid form within
the pressurized container can be tolerated so long as
the product that is dispensed contains the active
ingredient in solid form which is also solid (powder)
when the dispensed product strikes the skin.

10 For example, U.S. Patent No. 3,981,986 and
U.S. Patent No. 3,991,176 disclose antiperspirant
complexes which comprise a combination of a basic
aluminum-polyol compound, a zirconium compound and an
organic buffer. The complexes are said to be capable of
15 being used in conventional antiperspirant forms,
including aqueous solutions, aerosol sprays (including
powder-in-oil aerosol sprays) as well as creams,
lotions, and cream sticks. This patent further states
that the complexes can be formed in the container and
20 dispensed as a powder-in-oil aerosol spray wherein the
antiperspirant complex is a solid which is dispersed in
a non-solubilizing polar organic liquid. Thus, the
antiperspirant is in solid form within the dispenser and
remains in the solid form as discharged and applied to
25 the skin.

Similarly, U.S. Patent No. 3,288,681 discloses
an aerosol antiperspirant powder spray formed from a
dispersion containing an aluminum antiperspirant
compound, an alcohol, and a propellant. The product is
30 formulated by selecting the compounds and the relative
amounts of the compounds such that the antiperspirant is

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1 in the form of a solid in liquid dispersion within the
aerosol container and remains in insoluble form as it is
discharged from the container and applied to the skin.

U.S. Patent No. 3,876,758 discloses aerosol
5 dispensing systems for antiperspirants including an
antiperspirant component which is insoluble within the
container, and remains in solid form upon application to
the skin.

U.S. Patent No. 3,873,686 discloses an aerosol
10 antiperspirant formulation which, according to the
patent, was converted into a powder immediately on
leaving the aerosol container and landed on the human
skin in the form of a powder.

Some products use a volatile liquid combined
15 with product in a sealed container such as roll-on
deodorants or antiperspirants or the like. When the
product is applied from the container with a suitable
dispensing system, the volatile liquid portion of the
product evaporates leaving the desired active
20 ingredient.

It can thus be seen that formulation practice
with pressurized dispensing systems, especially systems
for dispensing difficultly soluble active ingredients,
has been to formulate the active ingredient into a
25 composition whose form within the dispenser is the same
as the form that the product is desired to exhibit
following discharge. That is, products which are
desired to be in liquid solution form upon application
are formulated within the pressurized container in
30 combination with propellant as solutions, and products
desired to be in the solid form upon application are

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1 formulated as a solid phase with propellant within the
container.

The present inventors have realized that
additional heretofore unprecedented freedom and
5 flexibility can be realized by adopting a new course in
the formulation of compositions to be dispensed from
active solids in dispersion in the container to active
ingredient in solution on application to the target.

The invention involves the formation of a
10 solution of one or more active ingredients in a suitable
solvent which is admixed with a volatile propellant or
volatile liquid. On addition of a propellant, or
volatile liquid, the propellant mixes with the solvent
causing one or more of the active ingredients to
15 precipitate in the homogeneous solvent-propellant
admixture. On dispensing the dispersed, at least
partially insoluble active ingredient in the propellant
and solvent mixture from a pressurized container
(aerosol can), the propellant quickly evaporates and the
20 previously insoluble active ingredient redissolves to
form a clear (or slightly cloudy initially) solution for
application. Alternatively, a volatile liquid
evaporates when a container is opened and its contents
applied, leaving the desired active ingredient in
25 soluble and activated form.

Disclosed are compositions and methods of
forming and applying the composition. Particularly
preferred are antiperspirant compositions solubilized in
alcohol and/or water and mixed with a volatile
30 propellant such as difluoroethane where the solvent and
propellant form a homogeneous mixture of suspended

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1 active ingredients within the pressure container but
when dispensed yield an essentially clear antiperspirant
solution of active ingredients without evidence of
solids on the skin.

5

More particularly, the present invention is directed to a composition capable of being dispensed from a container comprising a admixture of: at least one active ingredient; an amount of solvent effective to
10 dissolve said active ingredient at room temperature and atmospheric pressure; and a volatile propellant or liquid capable of forming a homogeneous mixture with said amount of solvent; at least a portion of said active ingredient being insoluble and dispersed in said
15 homogeneous mixture, whereby on dispensing the admixture, the admixture is capable of separating into volatile propellant or liquid and a solvent containing one or more active ingredients dissolved therein.

The present invention is further directed to a
20 method of preparing a product for dispensing from a container comprising admixing: at least one normally solid active ingredient; an amount of solvent effective to dissolve the active ingredient at room temperature and atmospheric pressure; and a volatile propellant or
25 liquid capable of forming a homogeneous mixture with said amount of solvent at least a portion of said active ingredient being insoluble and dispersed in said homogeneous mixture; said admixture effective on being dispensed to separate into volatile propellant; and a
30 solvent containing one or more active ingredients dissolved therein.

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1 The Figure is a cross-sectional schematic view
of a dispenser useful in the practice of the present
invention.

 The present invention is useful for dispensing
5 any of a wide variety of products. Examples include
products which may be applied as solutions to target
surfaces, such products including for example
antiperspirants, herbicides, pesticides, insect
repellents, coating compositions, adhesives, and the
10 like. Other examples include compositions which are
utilized as solutions which are simply sprayed into the
ambient atmosphere, without necessarily being directed
to a specific target surface. Examples of the latter
compositions include perfumes, room deodorizers,
15 fumigants, disinfectants, and the like. Other products
include the active ingredient in suitable solvent and a
volatile liquid which evaporates when the container is
opened.

 Without being bound by any particular
20 limitation as to form, the solutions upon discharge are
a solution of active ingredient completely dissolved in
a solvent for the active ingredient. As will be
described further herein below, the solution forms in
the ambient atmosphere essentially immediately upon
25 being discharged from the container. Thus, when the
composition is intended to be applied to a target
surface, it is preferably already in solution form when
it reaches that surface. In some embodiments, however,
it is permissible that the solution form upon impact
30 with the target surface. It should be recognized that

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1 compositions within the scope of this invention may
comprise more than one active ingredient of interest.

The dispensable composition is formed by
dissolving one or more active ingredients in a suitable
5 solvent which can comprise one or more components to
form a clear solution. Solutions approaching the
solubility limit of the actives in the solvent can be
used to provide maximum active ingredient in the
composition where needed. Lower amounts may be employed
10 depending on the requirements of the application. Once
the solution is prepared, the solution is contacted with
propellant under pressure or admixed with a volatile
liquid. The propellant is selected to form a
homogeneous mixture of solvent and propellant although
15 excess propellant may be employed. The volatile liquid
is any material forming at atmospheric pressure or above
a homogeneous admixture with the solvent wherein the
volatile liquid will evaporate rapidly when the
container is opened. The homogeneous mixture of solvent
20 and propellant or volatile liquid lowers the solubility
of active ingredient in the solvent causing the active
ingredient(s) to precipitate from the mixture as a
solid, often first appearing cloudy and then gelatinous
before heavy precipitation of solids occur. The amounts
25 of solvent and active ingredients and propellant or
volatile liquid are adjusted to produce a solid
dispersion of the active ingredient which is suspended
or can be suspended in the homogeneous solvent and
propellant or volatile liquid mixture by shaking the
30 mixture.

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1 On dispensing the dispersion formed previously
from a container, the propellant or volatile liquid
evaporates and the active ingredient re-dissolves to
form a solution which may be cloudy for an instant as
5 the solids re-dissolve in the solvent. Almost
immediately after evaporation of the propellant or
volatile liquid, the active ingredient is totally
dissolved in the solvent providing a solution which
forms a clear film on the target. Thus a solid
10 dispersion of "Chlorhydrol" or other active ingredient
in solvent and propellant under pressure can be applied
to the body or other target as a clear solution already
in effective form without unsightly powder present. On
dispersing the packaged dispersion of active ingredient
15 in solvent and volatile liquid, the volatile liquid
evaporates and the active ingredient re-dissolves in the
solvent during or shortly after application.

Any homogeneous solvent and propellant mixture
can be employed. We have found as propellants
20 non-chlorinated fluorocarbons, low molecular weight
ethers, hydrocarbons, such as lower alkanes, either
alone or mixed with each other or mixed with other
propellants produces excellent results allowing the
loading of high levels of active ingredient in the
25 mixture of solvent and propellant under pressure but
providing an essentially clear propellant free solution
of active ingredient for application at room temperature
and pressure.

The mixture can be prepared under pressure to
30 form the dispersion of active ingredient in a solvent
and propellant homogeneous mixture so long as the amount

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1 of active ingredient is maintained below the solubility
maximum of the active ingredient in the solvent at the
application conditions of temperature and atmospheric
pressure. The solution can be conveniently filled at
5 atmospheric conditions into aerosol cans in a
conventional manner. The clear solutions of active
ingredient in solvent can be filled with propellant.

One aspect of the present invention comprises
a system and method for dispensing an active ingredient,
10 the system comprising a sealed, valved container which
holds a gaseous propellant, the propellant comprising
one or more gas components, the container further
holding a dispersion of solid active ingredient slurried
in a solution of one or more of said gas components
15 dissolved in a liquid solvent for said active
ingredient, the amount of said active ingredient in said
dispersion not exceeding the solubility limit thereof in
said solvent, wherein the container is fitted with
externally actuatable valve means for dispensing said
20 dispersion from said container under pressure exerted by
said propellant wherein upon dispensing of said
dispersion said one or more dissolved gas components are
liberated therefrom and the dispensed solvent completely
dissolves the dispensed active ingredient. The method
25 is to establish a dispersion of a solid active
ingredient slurried in a solution of one or more of said
gas components dissolved in a liquid solvent for said
active ingredient, wherein the active ingredient is
present in said dispersion in an amount which does not
30 exceed the solubility limit thereof in said solvent, and
dispensing said dispersion from said container whereupon

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1 said one or more dissolved gas components evaporates
from the mixture allowing the dispensed solvent to
redissolve the active ingredient.

With reference to the Figure, dispensing
5 system 1 includes a container 2. Container 2 is a
sealed enclosure to which is fitted externally
actuatable dispensing means 3 for dispensing product
when desired from within container 2 to the ambient
atmosphere or to a target surface. Within container 2
10 are a dispersion 4 of active solids in a homogeneous
mixture of solvent and propellant 5.

Container 2 is preferably formed of metal, or
rigid plastic, which is inert to the contents of the
container. The container can be formed of one integral
15 piece, such as a drawn aluminum can, or it can be formed
in a conventional manner from several pieces including
pieces forming the sides and the upper shoulders to
which dispensing means 3 is fitted and a piece forming
the bottom which is fixed to the bottom edge of the side
20 of the container throughout the circumference thereof.
When desired, the interior of container 2 can be
provided with a coating to protect the container
material from corrosion or other adverse reactions with
the contents thereof. Suitable treatment can include
25 shellac or a thin polymeric barrier film applied to the
interior of container 2. Barrier packages may also be
used.

Dispensing means 3 includes a button 6, valve
means 7, and dip tube 8. Valve means 7 is preferably of
30 the type having a passage for product being dispensed,
and a valve operatively coupled to a spring means which

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1 urges the valve to a normally closed position preventing
passage of product therethrough except when the valve is
urged to an opening position by external actuation.
Button 6 is fitted to the discharge end of the passage
5 and contains an orifice appropriately configured to
permit passage therethrough of the solid and liquid
components of dispersion 4. The orifice of button 6 is
also appropriately dimensioned to provide the desired
spray pattern, including the desired angle through which
10 product is dispensed i.e. as a stream or spray, and the
desired droplet size, i.e., as a spray, a mist or a fog
of micro droplets.

Dip tube 8 is attached to the end of valve
means 7 within container 2 so as to be in fluid
15 communication with the discharge passage. Dip tube 8 is
a narrow, hollow tube dimensioned to convey the solid
and liquid components of dispersion 4. Preferably, the
lower end of dip tube 8 is at or near the bottom of
container 2, to maximize the amount of dispersion 4 that
20 can be discharged from the container before the contents
are considered to be fully spent.

Dispersion 4 comprises an intimate slurry of a
solid phase slurried in a homogeneous solvent and
propellant or volatile liquid mixture. The solid phase
25 includes the active ingredient to be dispensed. A
portion of the active ingredient may also be dissolved
in the solvent propellant mixture.

Active ingredients useful in the present
invention include solids which form solutions, in
30 solvents that are liquid at room temperature and at
atmospheric pressure. Any such solution should

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- 1 preferably contain dissolved therein a sufficient amount of the active ingredient such that the active ingredient is effective for its intended function. For instance, a solution of an antiperspirant would be desired to
- 5 contain about 2% to about 10% or more by weight, of antiperspirant upon application of the solution to the skin. The amount of any other active ingredient that should be present in a solution thereof, to be dispensed from the dispensing systems of the present invention,
- 10 will vary according to the identity of the active ingredient and its desired function. Such amounts can readily be determined from reference literature or from simple experimentation, for any particular active ingredient. The maximum amount of active ingredient
- 15 that can be dissolved in any particular solvent or solvent system is determined by the "solubility limit", by which term is meant the maximum amount of an ingredient that can be completely dissolved in a given volume of solvent in which no propellants or volatile
- 20 liquids are dissolved, at atmospheric pressure and the temperature at which the composition is to be dispensed.

Solvents useful in the dispensing systems of the present invention are generally characterized as being liquid at room temperature and pressure with

- 25 varying degree of volatility and are capable of solubilizing the desired active ingredient at room temperature and pressure and are also capable of dissolving a portion of the gaseous propellant or volatile liquid employed in the dispensing systems of
- 30 the present invention. Where that gaseous propellant is composed of more than one component, satisfactory

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- 1 solvents include those which can solubilize all such
propellant components to form a homogeneous propellant
and solvent mixture.

Suitable solvents for varying active

- 5 ingredients include water, mono and polyhydric alcohols
of two or more carbons such as lower alkanols, such as
ethanol, n-propyl alcohol, isopropyl alcohol, butanol,
polyols such as glycerine and propylene glycol,
dipropylene glycol, polyethylene glycols and the like;
10 esters, including lower alkyl esters of lower alcanoic
acids, such as ethyl acetate; ethers, such as diethyl
ether and methylethyl ether; alkanes; hydrocarbons,
kerosene, oils such as mineral or vegetable oils and
lower alkyl ketones, such as acetone and methylethyl
15 ketone. Satisfactory solvents can also include
one-phase mixtures of any of the foregoing. For
instance, satisfactory solvents for a system in
accordance with the present invention for dispensing
antiperspirant can comprise water, alcohol or a
20 water-alcohol mixture, depending upon the identity of a
particular compound or compounds used as the
antiperspirant active ingredient. The amount of solvent
is generally 20 to 60 wt. %, preferably 25 to 55 wt. %.

- The relative amounts of active ingredient,
25 solvent, and propellant are chosen to provide the
properties within the container and upon dispensing that
are desired as taught herein; identification of
effective amounts of each component is a straightforward
matter particularly with the guidance provided by the
30 Examples and Tables herein.

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- 1 Propellant system 5 is sealed within container
2. Sufficient propellant is present to exert a positive
pressure on dispersion 4 such that when valve means 3 is
actuated, an amount of dispersion 4 is forced from
5 within container 2 to the atmosphere.

Propellants useful in the practice of the
present invention can generally include any liquefiable
or compressible gaseous propellant conventionally used
in aerosol-type dispensers. For example, propellants
10 are selected from the group consisting of
non-chlorinated fluorocarbons, low boiling ethers, low
boiling hydrocarbons and mixtures thereof. Specific
examples included 1,1-difluoroethane, tetrafluoroethane,
and other non-chlorinated fluorocarbons, particularly
15 those with minor environmental consequence, propane,
isobutane, n-butane, dimethyl ether, and the like.

Suitable volatile liquids would be those
compounds which are liquid under conditions of use but
easily volatilized or evaporated under normal
20 atmospheric conditions or body heat or the like on
application from a container.

The propellant, or volatile liquid, single
compound or mixture, is selected to form a homogeneous
mixture with the active ingredient laden solvent usually
25 combined under pressure. At that point, solids
dissolved in the solvent precipitate forming a
dispersion of solid active ingredient in a homogeneous
mixture of propellant and solvent. Some active
ingredient may remain dissolved in the solvent and
30 perhaps in the propellant, or volatile liquid, depending
on solvent properties of active ingredient. Simple

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1 tests of selected solvent and propellant or volatile
liquid in glass pressure containers can be used to
determine homogeneous mixtures. Two phase mixtures,
while they would work with vigorous shaking prior to
5 application, are not preferred because the active
ingredient concentrates in one of the phases and may be
non-uniformly dispersed when the phases are not well
mixed. Excess propellant, in small quantity, may exist
as a separate phase for use in pressurizing the
10 container where necessary. The propellant solvent
mixture is selected, however, to give a homogeneous
mixture containing the dispersed active ingredient
solids under pressure in the aerosol container, and on
dispensing a solution of active ingredient and solvent
15 as the propellant vaporizes at atmospheric pressure.

In summary, the solvent making up the liquid
phase of dispersion 4 and the composition of the
propellant phase 5 are selected so that a portion of the
propellant is dissolved within the liquid phase of the
20 dispersion when the contents within container 2 are
fully pressurized. Distributing the propellant between
the gas phase 5 and the liquid phase of dispersion 4
displaces active ingredient from solution in the solvent
and assists in maintaining the desired slurry of solid
25 active ingredient in the solvent. The active ingredient
may be distributed between the solid phase of dispersion
4 and the solvent propellant forming the liquid phase of
dispersion 4. On dispensing the active ingredient re-
dissolves in the solvent as propellant is vaporized.

30 The dispensing system of the present invention
can be produced using techniques conventionally employed

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1 for manufacturing aerosol dispensing systems, with but
minor modifications. A solution is prepared of the
active ingredient dissolved in the solvent system
(comprising a single compound or a one-phase mixture of
5 compounds and/or solvents as the case may be). The
amount of active ingredient employed should not exceed
the solubility limit of that material in the solvent
system employed. In that way, the amount of active
ingredient present in the dispersion that subsequently
10 forms on addition of propellant --that is, present in
solution prior to addition of propellant or slurried in
solid form after addition of propellant and sealing the
can-- is also no greater than the solubility limit of
the active ingredient in the solvent when sprayed to
15 atmosphere or the target (propellant has evaporated).

An amount of solution containing solubilized
active ingredient is charged to the container 2. This
solution can be fed into the container using
conventional means. The propellant of choice for the
20 given system is fed into the container. The container
can be sealed before or after adding propellant.
Propellant can be fed using either through-the-valve or
under-the-cup filling techniques conventionally employed
in this industry for charging propellant gas to a
25 pressurized aerosol container. As the amount of
propellant present within the container increases, an
equilibrium portion thereof dissolves in the liquid
solvent present in the container. As the solubility of
the active ingredient in the solution of the propellant
30 and solvent decreases to become less than the solubility
of the active ingredient in the solvent per se,

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1 continued feeding of the propellant into the container
and solubilization thereof in the solvent causes the
active ingredient to form a precipitate which remains
slurried in the solvent. The propellant should not
5 interact with the active ingredient or the solvent as by
undergoing a chemical reaction or forming an
indispersible gel or other by-product. Determining
propellants that satisfy these criteria is a straight-
forward matter. In some cases the entire amount of
10 active ingredient will precipitate from the solvent,
whereas in other cases only a portion thereof will
precipitate such that the active ingredient remains
distributed between the solid phase and the solvent.

Sufficient propellant is charged to the
15 interior of the container 2 to precipitate active
ingredient from solution in the solvent and to establish
over the dispersion which is thus formed a sufficiently
high pressure such that when the dispensing means 3 is
actuated, the propellant drives the solid and solvent
20 components of the dispersion up dip tube 8 and out
through the dispensing means 3 to the atmosphere. The
final pressure of the fully charged dispenser is
generally about 20 to about 120 psig. The amount of
propellant in the container will generally be about 10%
25 or more by weight of all ingredients, preferably 25-90%,
and more preferably about 30 to about 70% by weight.

When the dispersion is thereafter discharged
from the container, the portion of the propellant gas
which had been dissolved in the solvent portion of the
30 dispersion evaporates into the atmosphere, and the solid
active ingredient that was dispensed rapidly

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1 resolubilizes into the solvent portion that had been
discharged from the container. As a result, the
dispensing system of the present invention effectively
provides a solution of active ingredient in a solvent
5 therefor to the atmosphere or to any particular desired
target surface. This avoids the normal heavy spray of
solids when dispensing a dispersion of solids from the
aerosol can. If desired, preformed mixtures of active
ingredient, solvent, and propellant can be prepared and
10 the aerosol container 2 filled with the mixture under
pressure. If desired, solid active ingredient can be
added to solvent and propellant to form such mixtures
for filling provided the active ingredient is formulated
to dissolve in the solvent when propellant vaporizes
15 from the mixture during use.

Products can also be prepared from active
ingredient dissolved or combined with a solvent
therefore, and a volatile liquid which can be packaged
conventionally in pressure and non-pressure packages.
20 On dispensing the product from the container, such as a
liquid antiperspirant, the volatile liquid evaporates
and the active ingredient re-dissolves in the solvent
thereby being more available than a powder version of
the same active ingredient.

25 Applying the active ingredient as a solution
affords a number of advantages including even and
thorough application; and the ability to adhere to
surfaces to which solids would not readily adhere. In
addition, for products whose appearance upon application
30 is a significant factor, application in the form of a
solution is preferable because the solution is clear and

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1 leaves no unsightly residue whereas application of a
powder will create a visible residue or film.
Antiperspirants are a significant example of such a
product whose appearance upon discharge is an important
5 factor affecting the attractiveness and salability of
the product. In addition, applying the active
ingredient as a solution avoids the necessity of
dissolving the active ingredient after it has been
dispensed as occurs when antiperspirants (as in powder-
10 in-oil aerosols) must be activated and solubilized by
body moisture. In addition, faster drying formulas with
less irritating spray particles can be developed.

Another significant advantage of the invention
described herein is that it permits the dispensing of a
15 solution containing active ingredient in higher amounts
than previously achieved. The system holds more active
ingredient than can be kept in solution within the
pressurized container, yet manages to dispense that
active ingredient in a completely solubilized form.

20 The invention permits tailoring the amounts of
each component, particularly the propellant, to achieve
desirable results unconstrained by the solubility of the
active ingredient in the homogeneous solution formed by
the propellant in the solvent. Increasing the amount of
25 propellant --which is now permitted by the present
invention, even as the active ingredient precipitates--
permits establishing better atomization and quicker
drying of the dispensed solution. The invention also
permits using less propellant per amount of active
30 ingredient.

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1 The present invention also permits the
formulation of dispensing systems using propellants that
are environmentally benign, without being unduly limited
to systems that remain in solution form in the
5 pressurized container.

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EXAMPLE 1

Pressurized dispensing systems for antiperspirants were prepared containing the components listed in the following Table 1, in the amounts shown in the table. Solutions were prepared containing all the components except the propellant. The solution was sealed in valved glass bottles, and then the indicated amount of propellant was injected through the valve.

A powder was seen to precipitate upon addition of the propellant. The powder was a fine white dispersion that dispersed easily upon gentle shaking, and thereafter settled slowly. When samples of these preparations were sprayed, under pressure from the propellant, the spray did not contain noticeable solid particles and rapidly formed a solution which was clear on the skin and rapidly dried to an unnoticeable film.

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TABLE 1

AMOUNTS (wt. %)

[illegible]

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EXAMPLE 2

Pressurized dispensing systems for antiperspirants were prepared containing the components listed in the following Tables 2, 2A, and 2B, in the amounts shown. The solution was sealed in valved glass bottles and the amount of propellant injected through the valve. Various amounts of active ingredient indicate the advantage of forming a precipitate within the can or jar which is soluble in solvent on dispensing, in that more active ingredient can be dispensed and the amount of propellant used per unit of active ingredient can be reduced. There is some change in the amount of active ingredient which is soluble in the solvent propellant solution in the pressure container over time.

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TABLE #2
Effect of Reach 501 Concentration with Various Propellants
Amount - % by weight

Example	1	2	3	4	5	6	7	8	9	10	11	12
Reach 501 (50% Aqueous Solution)	15.0	12.5	10.0	7.5	15.0	12.5	10.0	7.5	15.0	12.5	10.0	7.5
SD#40 Anhyd. (100% ethanol)	45.0	47.5	50.0	52.5	45.0	47.5	50.0	52.5	45.0	47.5	50.0	52.5
P152A 1,1-Difluoroethane)	40.0	40.0	40.0	40.0	-	-	-	-	-	-	-	-
P-152A/A31 75% 1,1- Difluoroethane 25% Isobutane	-	-	-	-	-	-	-	-	40.0	40.0	40.0	40.0
DME/A31 60% Dimethyl ether 40% Isobutane	-	-	-	-	40.0	40.0	40.0	40.0	-	-	-	-
% Propellant Product Still Clear	36.0	37.2	39.7	40.0	26.4	31.8	36.7	39.5	31.0	31.8	36.6	40.0
% Propellant @ Approx. PPT. Point	37.6	37.8	40.0	-	28.2	34.9	37.8	-	32.0	32.8	37.9	-
Finish Product All Propellant Added	Powder Susp.	Milky Gel	Milky	Hazy Soln.	Powder Susp.	Powder Susp.	Milky	Milky Sl. Gel	Powder Susp.	Powder Susp.	Milky	Hazy Soln.
Day 1 @ R.T. Day 10 @ R.T.	Powder Susp.	Gels Thins @ Shake	Gels Thins @ Shake	Milky Soln.	Powder Susp.	Powder Susp.	Milky @ Gel Thins	Gels Thins @ Shake	Powder Susp.	Powder Sl. Gel Thins @ Shake	Powder Sl. Gel Thins @ Shake	Gels Thins @ Shake

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TABLE 2A

EFFECT OF VARYING CHLORHYDROL AND PROPELLANT CONTENTS
(all amounts in wt.%)

Test No.	1	2	3	4	5	6	7	8
Chlorhydrol (50% Aq. Solution)	15.0	12.5	10.0	7.5	20.0	17.5	15.0	12.5
SD # 40 (100% ethanol)	45.0	47.5	50.0	52.5	40.0	42.5	45.0	37.5
PI52A (1,1-difluoroethane)	40.0	40.0	40.0	40.0	40.0	40.0	40.0	50.0
Max. % Propellant w/ Product Still Clear	25	34	40	40	---	---	---	---
% Propellant at Start of Precip.	18.5	36.2	---	---	28.8	32.5	---	---
Final Product Appearance Held at Room Temperature at :	Powder susp.	Milky Starts to gel	Hazy No ppt.	Clear	Powder susp.	Powder susp.	Powder susp.	Powder susp.
day 3	---	---	---	---	Powder susp.	Powder susp.	---	---
day 5	---	---	---	---	---	---	Powder susp.	Powder susp.
day 10	Powder susp.	Gel	Gel thins with shaking	Slight hazy solut.	---	---	---	---

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TABLE 2A - CONTINUED

EFFECT OF VARYING CHLORHYDROL AND PROPELLANT CONTENTS
(all amounts in wt. %)

Test No.	9	10	11	12	13	14	15	16
Chlorhydrol (50% Aq. Solution)	10.0	5.0	2.5	17.5	14.6	11.7	5.8	2.9
SD # 40 (100% ethanol)	30.0	15.0	7.5	42.5	35.4	28.3	14.2	7.1
PI52A (1,1-difluoroethane)	60.0	80.0	90.0	40.0	50.0	60.0	80.0	90.0
Max. % Propellant w/ Product Still Clear	---	---	---	---	---	---	---	---
% Propellant at Start of Precip.	---	---	---	---	---	---	---	---
Final Product Appearance Held at Room Temperature at :	Powder susp.	Powder susp.	Powder susp.	Powder susp.	Powder susp.	Powder susp.	Powder caked to bottom	Powder caked to bottom
day 1	---	---	---	---	---	---	---	---
day 3	---	---	---	---	---	---	---	---
day 5	Powder susp.	Powder is susp. and caked on bottom	Powder is susp. and caked on bottom	Powder susp.	Powder susp.	Powder susp.	same as above	same as above
day 10	---	---	---	---	---	---	---	---

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TABLE 2B
EFFECT OF VARYING CHLORHYDROL CONTENTS
(all amounts in wt. %)

Test No.	1	2	3	4	5	6	7	8	9	10	11	12
Chlorhydrol (50% aq. Solution)	15.0	12.5	10.0	7.5	17.5	20.5	15.0	12.5	10.0	7.5	20.0	17.5
SD # 40 Ethanol	45.0	47.5	50.0	52.5	42.5	40.0	45.0	47.5	50.0	52.5	40.0	42.5
P152A/A31: 75%-1, 1- difluoroethane 25%-isobutane	40.0	40.0	40.0	40.0	40.0	40.0	---	---	---	---	---	---
DME/A31: 60%-dimethylether 40%-isobutane	---	---	---	---	---	---	40.0	40.0	40.0	40.0	40.0	40.0
Max. % Propellant w/ Product Still Clear	28.0	30.0	33.3	40.0	---	---	28.4	31.5	31.9	40.0	---	---
% Propellant at Start of Precip.	31.2	31.5	34.6	---	---	---	29.4	33.0	35.3	---	---	---
Final Product Appearance Held at Room Temperature at: day 1	Powder susp.	Powder susp.	Powder susp.	Hazy soln.	Powder susp.	Large par- ticles settling	Powder susp.	Powder, slight gel	Milky, starts to gel	Hazy starts to gel	Powder susp.	Powder susp.
day 3	---	---	---	---	---	---	---	---	---	---	Powder susp.	Powder susp.
day 7	---	---	---	---	Powder susp.	Colled to bottom	---	---	---	---	---	---
day 10	Powder susp.	Powder susp.	Powder susp.	Milky Powder	---	---	Powder susp.	Powder with gel, thins with shaking	Powder with gel, thins with shaking	Gel, thins with shaking	---	---

SUBSTITUTE SHEET (RULE 26)

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1 WHAT IS CLAIMED IS:

1. A composition capable of being dispensed from a container comprising a admixture of:
 - at least one active ingredient;
 - 5 an amount of solvent effective to dissolve said active ingredient at room temperature and atmospheric pressure; and
 - a volatile propellant or liquid capable of forming a homogeneous mixture with said amount of
 - 10 solvent;
- at least a portion of said active ingredient being insoluble and dispersed in said homogeneous mixture, whereby on dispensing the admixture, the admixture is capable of separating into volatile
- 15 propellant or liquid and a solvent containing one or more active ingredients dissolved therein.
2. A composition according to Claim 1 wherein said active ingredient is an antiperspirant.
3. A composition according to Claims 1 or 2
- 20 wherein the solvent is selected from the group consisting of water, lower alcohols, glycols, esters, ethers, ketones and mixture thereof.
4. A composition according to any of Claims 1-3 wherein the propellant is selected from the group
- 25 consisting of non-chlorinated fluorocarbons, low boiling ethers, low boiling alkanes, and mixtures thereof.
5. A composition according to any of Claims 1-4 wherein the propellant is selected from the group consisting of normal butane, propane, difluoroethane,
- 30 tetrafluoroethane, isobutane, dimethyl ether and mixtures thereof.

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- 1 6. A composition according to any of Claims
1-5 wherein said propellant comprises at least about 10%
or more of the homogeneous mixture.
- 5 7. A composition according to any of Claims
1-6 where the propellant is selected from the group
consisting of non-chlorinated fluorocarbons, low boiling
ethers, low boiling alkanes, and mixtures thereof.
- 10 8. The composition according to any of Claims
1-7 wherein the solvent is selected from the group
consisting of water, lower alcohols, polyols, esters,
ethers, ketones and mixtures thereof.
- 15 9. A method of preparing a product for
dispensing from a container comprising admixing:
at least one normally solid active ingredient;
an amount of solvent effective to dissolve the
active ingredient at room temperature and atmospheric
pressure;
and a volatile propellant or liquid capable of
forming a homogeneous mixture with said amount of
20 solvent at least a portion of said active ingredient
being insoluble and dispersed in said homogeneous
mixture;
said admixture effective on being dispensed to
separate into volatile propellant;
25 and a solvent containing one or more active
ingredients dissolved therein.
10. The method of Claim 9 wherein said active
ingredient is an antiperspirant.
11. The method of Claims 9 or 10 wherein said
30 solvent is selected from the group consisting of water,

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1 lower alcohols, glycols, esters, ethers, ketones and mixtures thereof.

12. The method of any of Claims 9-11 wherein the propellant is selected from the group consisting of
5 non-chlorinated fluorocarbons, low boiling ethers, low boiling alkanes, and mixtures thereof.

13. The method of any of Claims 9-12 wherein the propellant is selected from the group consisting of difluoroethane, tetrafluoroethane, isobutane, normal
10 butane, propane, dimethyl ether and mixtures thereof.

14. A system for dispensing an active ingredient, the system comprising a sealed, valved container and within the container a gaseous propellant, the propellant comprising one or more gas components,
15 the container further holding a dispersion comprising solid active ingredient slurried in a solution of one or more of said gas components dissolved in a liquid solvent for said active ingredient, the amount of said active ingredient in said dispersion not exceeding the
20 solubility limit thereof in said solvent, and externally actuatable valve means fitted to said container for dispensing said dispersion from said container under pressure exerted by said propellant wherein upon dispensing of said dispersion said one or more dissolved
25 gas components are liberated therefrom and the dispensed solvent completely redissolves the dispensed active ingredient.

15. A system according to Claim 14 wherein said solvent is selected from the group consisting of
30 water, lower alcohols, glycols, esters, ethers, ketones and mixtures thereof.

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1 16. A system according to Claims 14 or 15
wherein said propellant is selected from the group
consisting of non-chlorinated fluorocarbons, low boiling
ethers, low boiling alkanes, and mixtures thereof.

5 17. A system according to any of Claims 14-16
wherein said active ingredient is an antiperspirant.

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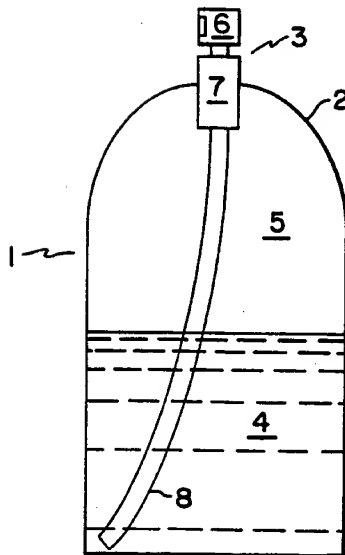
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1 / 1

Figure



INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/07513

A. CLASSIFICATION OF SUBJECT MATTER																				
IPC(S) : A61K 9/08; 7/32 US CL : 424/401																				
According to International Patent Classification (IPC) or to both national classification and IPC																				
B. FIELDS SEARCHED																				
Minimum documentation searched (classification system followed by classification symbols)																				
U.S. : 424/401, 46, 65, DIG 1; 252/305; 222/402.1																				
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched																				
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)																				
C. DOCUMENTS CONSIDERED TO BE RELEVANT																				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.																		
Y	US, A, 5,118,494 (SCHULTZ ET AL) 02 June 1992; see entire document.	1-3, 9-11, 14-16																		
Y	Research & Development, "Fluorocarbon and Dimethyl Ether Aerosol Propellants", (STERLING), December 1982, pages 50-52.	1-3, 9-11, 14-16																		
Y	US, A, 3,981,986 (RUBINO) 21 September 1976; see entire document.	1-3, 9-11, 14-16																		
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.																				
<table border="0"> <tr> <td>* Special categories of cited documents:</td> <td>T</td> <td>later documents published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</td> </tr> <tr> <td>*A* document defining the general state of the art which is not considered to be part of particular relevance</td> <td>X</td> <td>document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</td> </tr> <tr> <td>*E* earlier document published on or after the international filing date</td> <td>Y</td> <td>document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, each combination being obvious to a person skilled in the art</td> </tr> <tr> <td>*L* document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</td> <td>*A*</td> <td>document member of the same patent family</td> </tr> <tr> <td>*O* document referring to an oral disclosure, use, exhibition or other means</td> <td></td> <td></td> </tr> <tr> <td>*P* document published prior to the international filing date but later than the priority date claimed</td> <td></td> <td></td> </tr> </table>			* Special categories of cited documents:	T	later documents published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	*A* document defining the general state of the art which is not considered to be part of particular relevance	X	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone	*E* earlier document published on or after the international filing date	Y	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, each combination being obvious to a person skilled in the art	*L* document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A*	document member of the same patent family	*O* document referring to an oral disclosure, use, exhibition or other means			*P* document published prior to the international filing date but later than the priority date claimed		
* Special categories of cited documents:	T	later documents published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention																		
A document defining the general state of the art which is not considered to be part of particular relevance	X	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone																		
E earlier document published on or after the international filing date	Y	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, each combination being obvious to a person skilled in the art																		
L document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A*	document member of the same patent family																		
O document referring to an oral disclosure, use, exhibition or other means																				
P document published prior to the international filing date but later than the priority date claimed																				
Date of the actual completion of the international search 25 AUGUST 1994		Date of mailing of the international search report 08 NOV 1994																		
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230		Authorized officer RAJ BAWA, P.H.D. Telephone No. (703) 308-2351																		

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US94/07513

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claims Nos.: 4-8, 12, 13 & 17
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐

The additional search fees were accompanied by the applicant's protest.

☐

No protest accompanied the payment of additional search fees.